

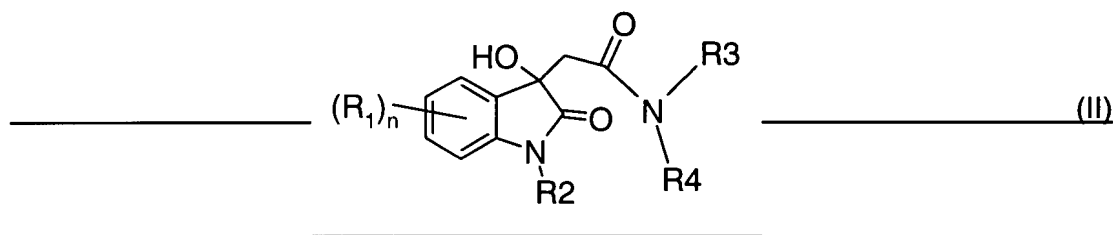
## IN THE CLAIMS

Kindly replace the prior claims listing by the following listing:

1-17: (cancelled).

18. (currently amended): A method for the manufacture of pharmaceuticals or of a compound of the formula II defined below,

comprising a method for the manufacture of amides of the formula II,



wherein n is a number from 0 to 4,

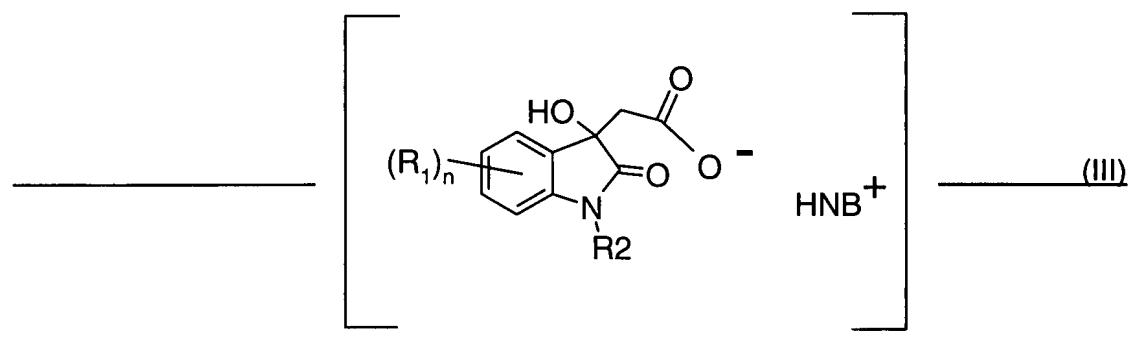
each R<sub>1</sub> is, independently of the other substituents R<sub>1</sub>, unsubstituted or substituted alkyl, unsubstituted or substituted aryl, unsubstituted or substituted heterocyclyl, alkylsulfonyl, sulfonyl alkyl, N-mono- or N,N-disubstituted or unsubstituted aminosulfonyl alkyl, hydroxy, mercapto, nitro, halogen, cyano, carboxamido, N-mono- or N,N-disubstituted carboxamido, carboxhydrazido, unsubstituted or substituted alkoxy carbonyl, unsubstituted or substituted alkoxy, formyl or other alkanoyl, unsubstituted or substituted alkenyl, unsubstituted or substituted alkynyl, unsubstituted or substituted cycloalkyl, alkanoyloxy, N-mono- or N,N-disubstituted or unsubstituted amino, unsubstituted or substituted hydrazino, or is a residue of a boronic acid or an ester thereof;

R<sub>2</sub> is hydrogen or unsubstituted or substituted alkyl, unsubstituted or substituted alkoxy carbonyl, unsubstituted or substituted arylsulfonyl, unsubstituted or substituted alkylsulfonyl, unsubstituted or substituted aryl, carbamoyl or N-mono- or N,N-disubstituted carbamoyl, silyl substituted by three moieties independently selected from unsubstituted or substituted alkyl and substituted or unsubstituted aryl, or acyl, and

R' is unsubstituted or substituted alkyl,

and R3 and R4 are, independently of each other, unsubstituted or substituted alkyl or together form an unsubstituted or substituted alkylene bridge (thus forming a ring with the binding nitrogen) or an alkylene bridge to which a phenyl or a C<sub>3</sub>-C<sub>8</sub>-cycloalkyl ring is condensed at two vicinal carbon atoms of the alkylene bridge

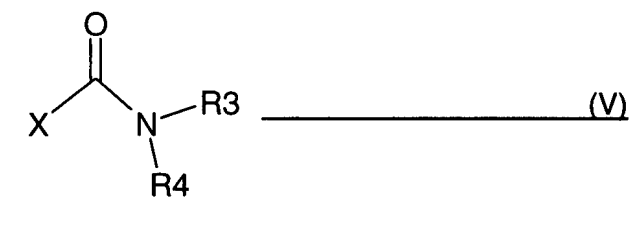
where a starting material of the formula III,



wherein n, R<sub>1</sub> and R<sub>2</sub> have the meanings given under formula II and NB is a tertiary nitrogen base where the nitrogen is not part of a ring,

is reacted

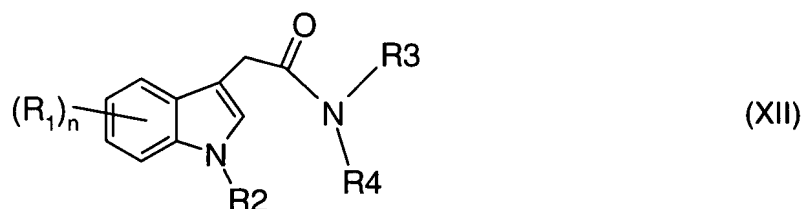
(b) with an active amido carbonic acid derivative of the formula V,



wherein X is halogen and R3 and R4 are as defined under formula II, to give the corresponding compound of the formula II,

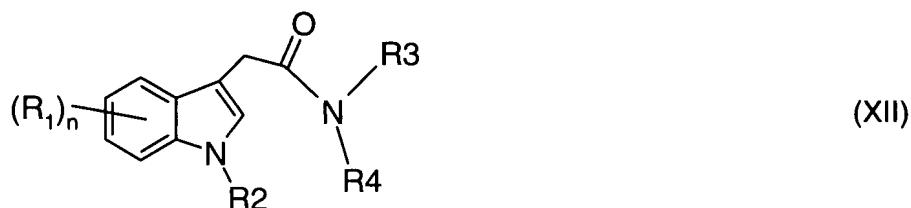
~~A method according to claim 1, and further comprising reducing the indolone moiety in [[a]] compound of the formula II wherein n, R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> are, independently of each other, as defined in claim 1, and R<sub>1</sub> is unsubstituted or substituted alkyl, unsubstituted or substituted aryl, unsubstituted or substituted heterocyclyl, alkylsulfonyl, sulfonyl alkyl, N-mono or N,N-disubstituted or unsubstituted aminosulfonyl alkyl, hydroxy, mercapto, nitro, halogen, cyano, carboxamide, N-mono or N,N-disubstituted carboxamide, unsubstituted or substituted alkoxy carbonyl, unsubstituted or substituted alkoxy, formyl or other alkanoyl, unsubstituted or substituted alkenyl, unsubstituted or substituted alkynyl, unsubstituted or substituted cycloalkyl, alkanoyloxy, N-mono or N,N-disubstituted or unsubstituted amino, or is a residue of a boronic acid or an ester thereof, in the presence of a complex hydride.~~

19. (previously presented): The method according to claim 18 wherein as reductant a borane di-lower alkyl sulfide is used, resulting in the formation of the corresponding indole of the formula XII



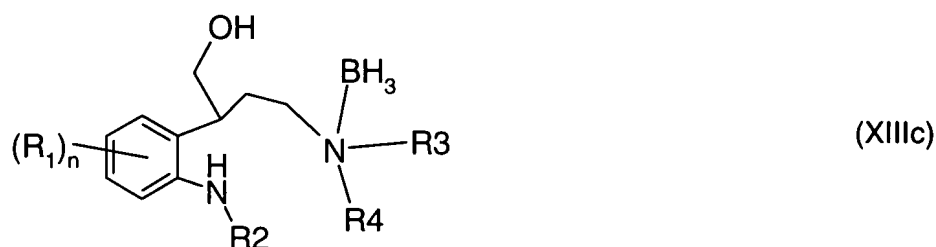
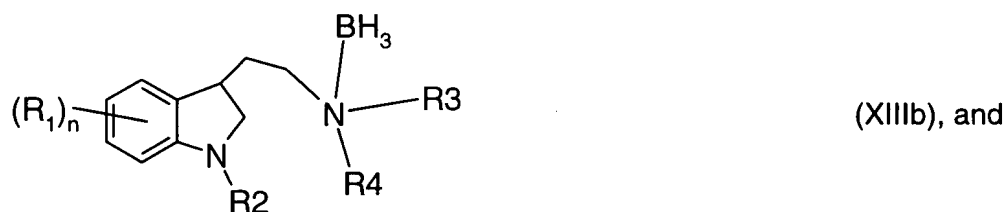
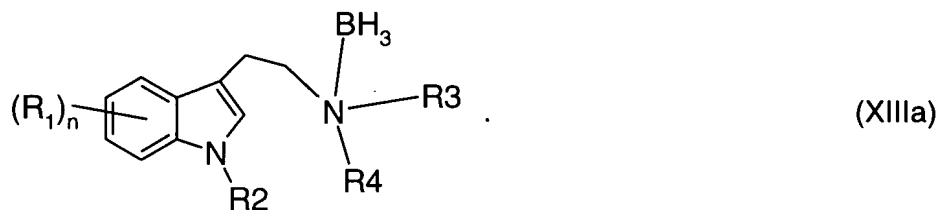
wherein the symbols and moieties are as defined in claim 18.

20. (currently amended): A compound of the formula XII



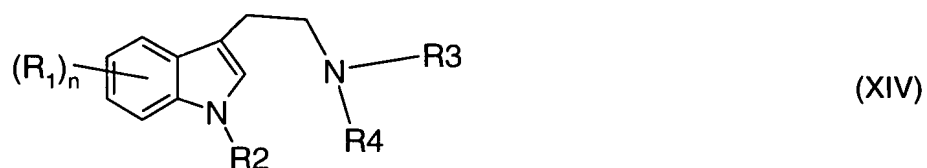
wherein n, R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> are as defined for formula II in claim 18, provided that R<sub>1</sub> is not alkoxy which is unsubstituted or substituted 5-methoxy if n is 1.

21. (previously presented): The method according to claim 18 where reaction of the compound of the formula II takes place in the presence of an alkali metal borohydride and a boron trifluoride etherate, yielding a mixture containing compounds of the formulae XIIIa, XIIIb and XIIIc,



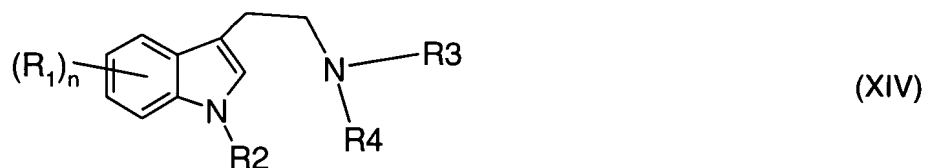
wherein n, R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> are as defined in claim 18 for the starting compounds of the formula II.

22. (original): A process according to claim 21, further comprising the conversion of the mixture of compounds XIIIa, XIIIb and XIIIc into a compound of the formula XIV



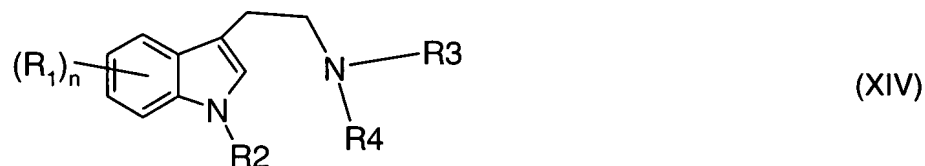
wherein n, R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> are as defined under formula XIIIa, XIIIb and XIIIc in claim 21, by reaction with diazabicyclo[2.2.2]octane and subsequent dehydrogenation or oxidation with an oxidant.

23. (currently amended): A compound of the formula XIV

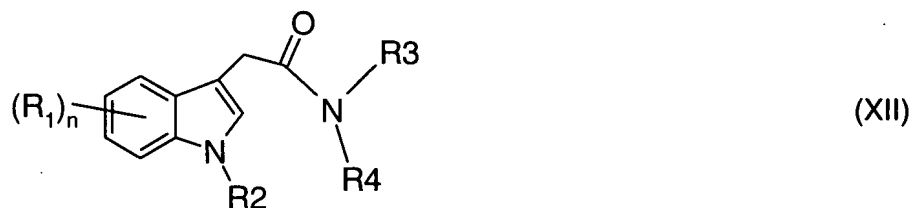


wherein n is a number from 1 to 4, R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> are as defined in claim [[1]]18 and R<sub>1</sub> is a residue of a boronic acid or ester thereof, lower alkyl, lower alkyl substituted by up to three moieties selected from N,N-di-lower acylamino and N-lower acylamino, C<sub>3</sub>-C<sub>10</sub>-cycloalkyl, C<sub>3</sub>-C<sub>4</sub>alkoxy~~C<sub>2</sub>-C<sub>4</sub>alkoxy~~, nitro, halogen, lower alkanoyloxy, unsubstituted or substituted aryl, unsubstituted or lower alkyl substituted and/or mono- or di-oxosubstituted nitrogen-heterocyclenyl or nitrogen-heterocyclyl, sulfonyl alkyl, mercapto, C<sub>2</sub>-C<sub>8</sub>alkanoyl, unsubstituted or substituted alkenyl, or unsubstituted or substituted alkynyl, or a salt thereof.

24. (currently amended): A method according to claim 22, wherein the~~Conversion of a compound of the formula XIV according to claim 22-~~



or of the formula XII,



where n, R<sub>1</sub>, R<sub>3</sub> and R<sub>4</sub> are as defined in claim 22 and R<sub>2</sub> is hydrogen, respectively, further is converted by introduction of a moiety R<sub>2</sub> which is unsubstituted or substituted alkyl, unsubstituted or substituted alkoxy carbonyl, unsubstituted or substituted aryl, carbamoyl, N-mono- or N,N-disubstituted carbamoyl, silyl substituted by three moieties independently selected from unsubstituted or substituted alkyl and substituted or unsubstituted aryl, or acyl; wherein unsubstituted or substituted alkyl is introduced by reaction with a strong base with a corresponding unsubstituted or substituted alkyl derivative of the formula XV,

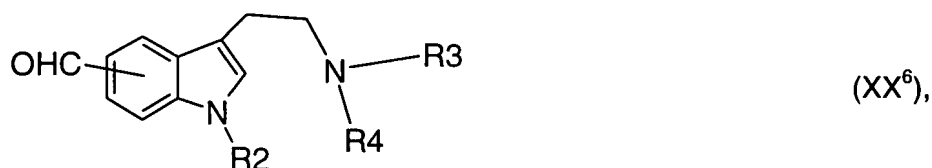
Alk-L

(XV)

wherein Alk is unsubstituted or substituted alkyl, unsubstituted or substituted alkoxycarbonyl, unsubstituted or substituted aryl, carbamoyl, N-mono- or N,N-disubstituted carbamoyl, and L is a leaving group, to give the corresponding compound of the formula XII or XIV wherein R<sub>2</sub> is unsubstituted or substituted alkyl; or acyl is introduced by reaction with the corresponding acylhalogenides or mixed or symmetrical acid anhydrides with one or two of the corresponding acyl moieties; or the silyl derivatives are introduced using the corresponding silylhalogenides, respectively; or a method for the synthesis of a tryptamine derivative ~~having pharmacologically useful properties~~ comprising said process~~conversion~~.

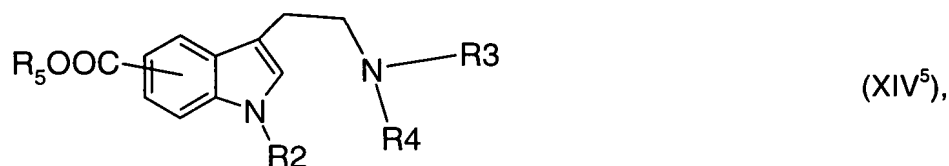
25-30. (cancelled).

31. (currently amended): A process for the reaction of a compound of the formula XIV as defined in claim 22 where n is 1 and R<sub>1</sub> is halogen, comprising converting it into the corresponding compound of the formula XX<sup>6</sup>,

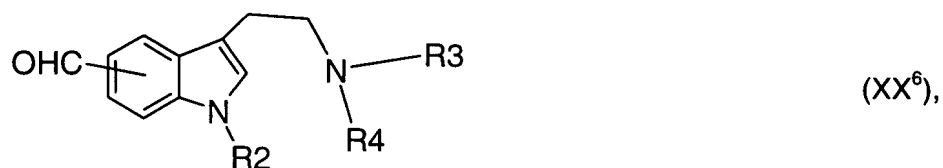


wherein R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> are as defined for the compound of the formula XIV, by reaction with first a lithium alkyl compound to form the lithio derivative and then with DMF or triethyl formate, to obtain the compound of the formula XX<sup>6</sup> after hydrolysis; or a method for the synthesis of a tryptamine derivative ~~having pharmacologically useful properties~~ comprising said process.

32. (currently amended): A compound of the formula XIV<sup>5</sup>

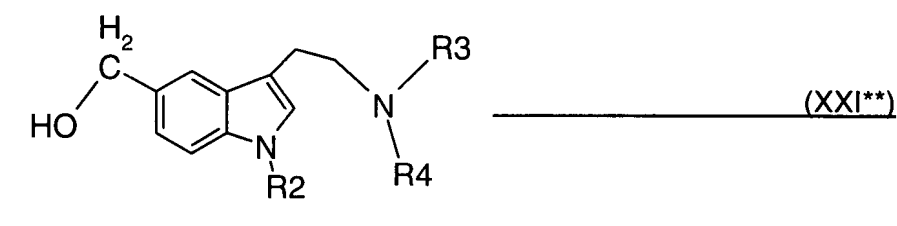
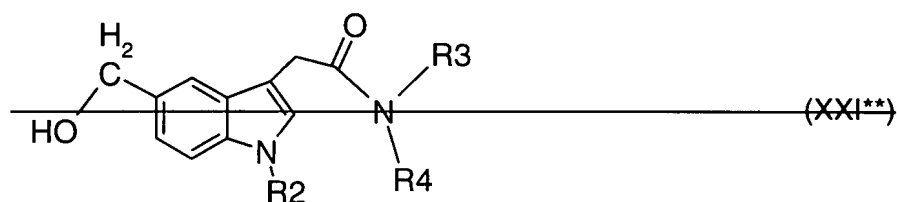


or of the formula XX<sup>6</sup>



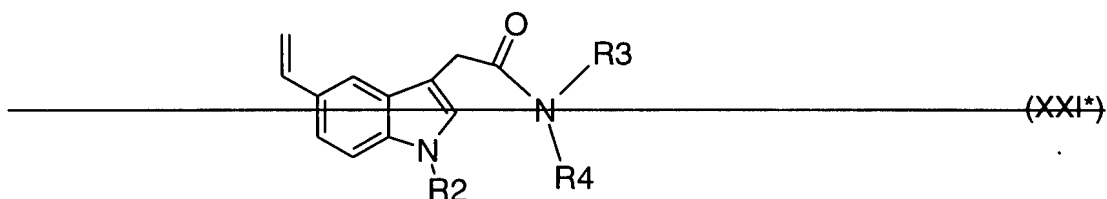
wherein R2, R3, ~~and R4 and R<sub>6</sub>~~ are as defined in claim ~~[[1]]18~~ for formula II, provided that one of R3 or R4 is not methyl and R3 and R4 together are not phthalyl, and R<sub>5</sub> is unsubstituted or substituted alkyl or unsubstituted or substituted aryl, or a salt thereof.

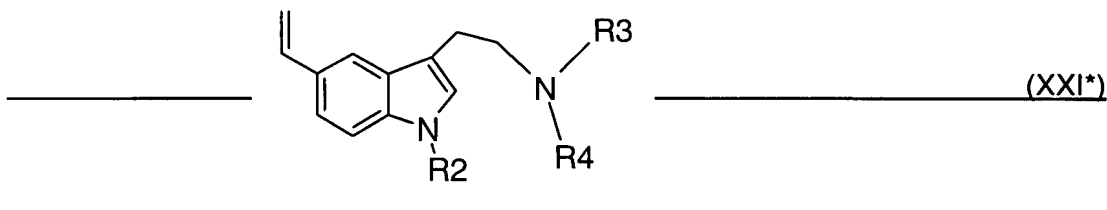
33. (currently amended): A process for the manufacture of a compound of the formula XXI\*\*



wherein R2, R3 and R4 have the meanings indicated for compounds of the formula XX<sup>6</sup> in claim 31, by reduction of the aldehyde carbonyl in the compound of ~~the~~ formula XX<sup>6</sup> in the presence of a selective transition metal catalyst; or a method for the synthesis of a tryptamine derivative ~~having pharmacologically useful properties~~ comprising said process.

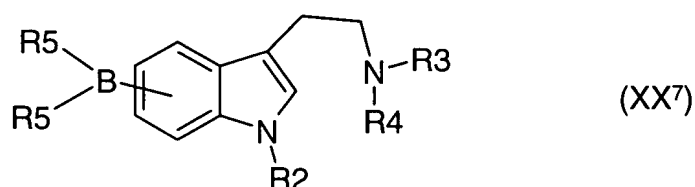
34. (currently amended): A process for the manufacture of a compound of the formula XXI\*,





wherein R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> have the meanings indicated for compounds of the formula XX<sup>6</sup> in claim 31, by conversion of a compound of the formula XX<sup>6</sup> as defined in claim 31 into the corresponding compound of the formula XXI\* by reaction with a Wittig or Wittig Horner reagent in the presence of a suitable base; or a method for the synthesis of a tryptamine derivative ~~having pharmacologically useful properties~~ comprising said process.

35. (currently amended): A process for the reaction of a compound of the formula XIV as defined in claim 22 where n is 1 and R<sub>1</sub> is halogen, comprising converting it into the corresponding compound of the formulae XX<sup>7</sup>,



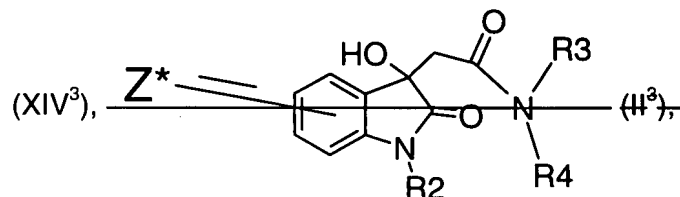
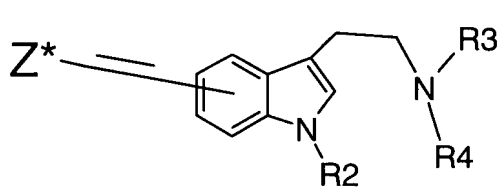
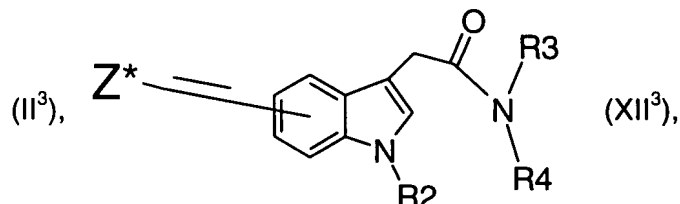
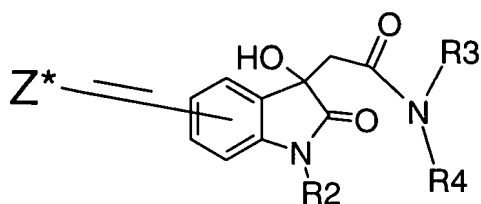
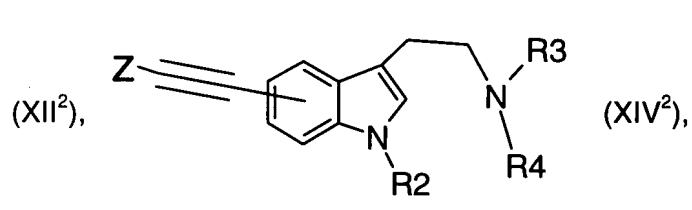
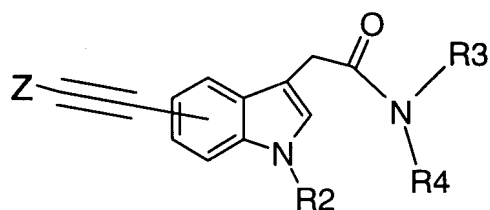
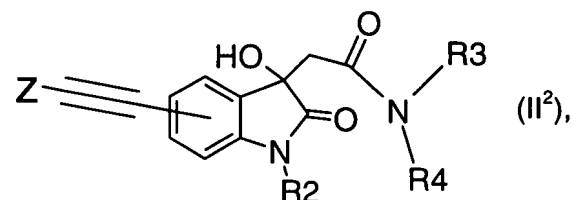
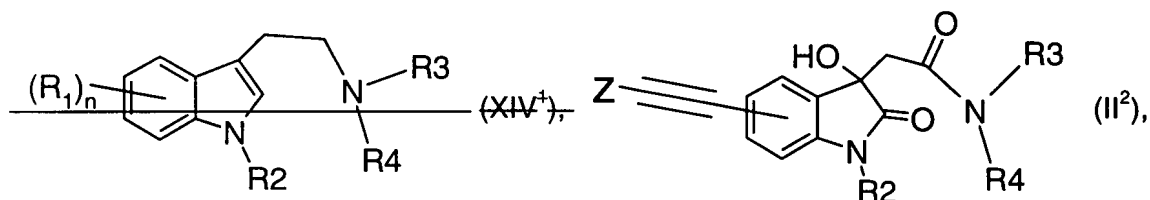
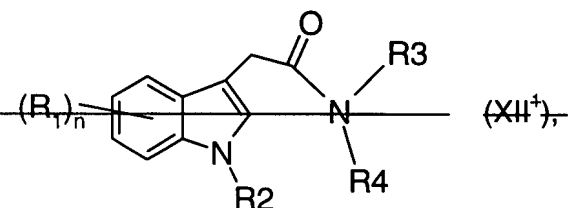
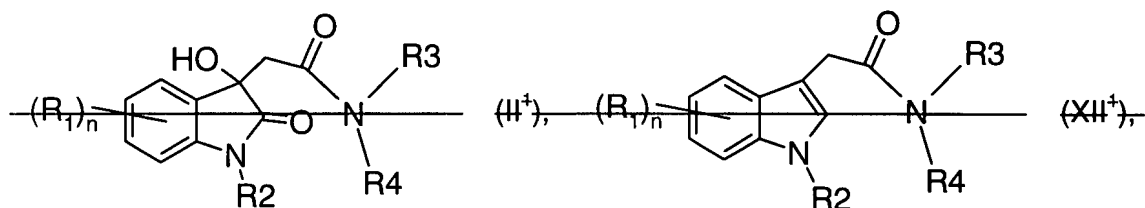
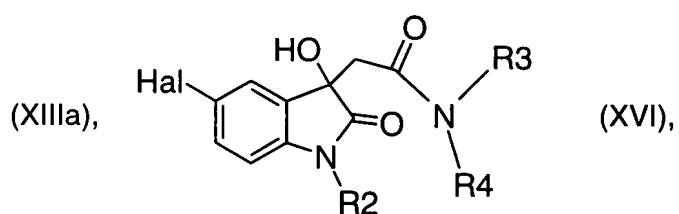
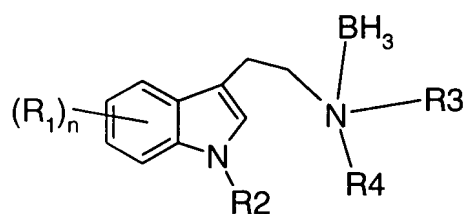
wherein R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> are as defined for the compound of the formula XIV, and each of R<sub>5</sub> independently is hydroxy or an alkoxy residue of a lower alcohol, or the 2 residues R<sub>5</sub> together are C<sub>2</sub>-C<sub>8</sub>alkylene-dioxy, by reaction with first a lithium alkyl compound to form the lithio derivative, and then with an ester of boric acid B,

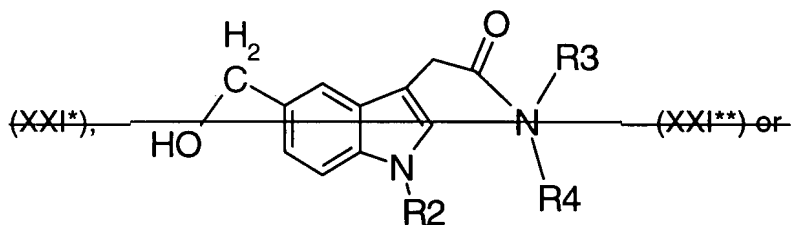
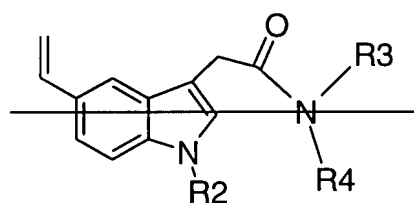
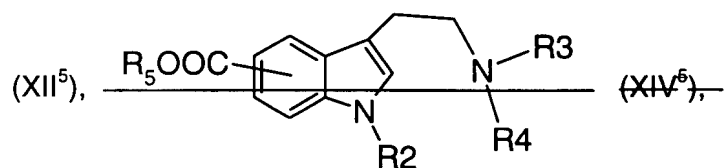
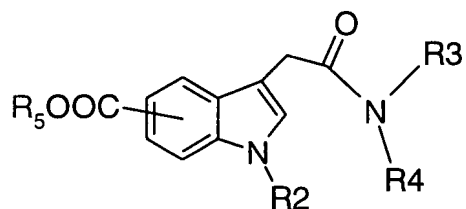
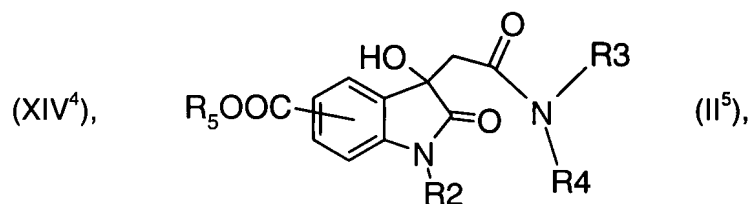
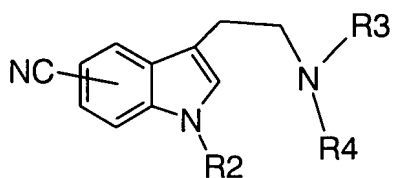
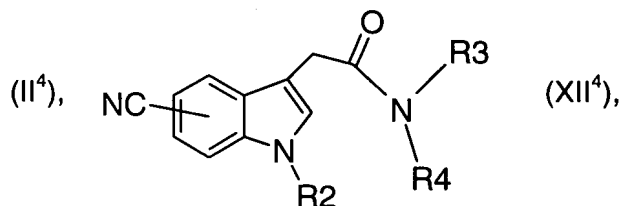
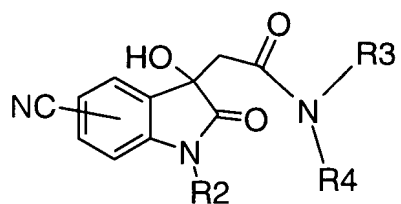
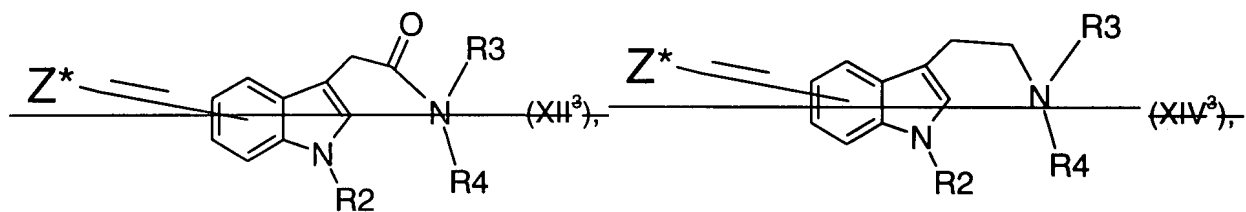


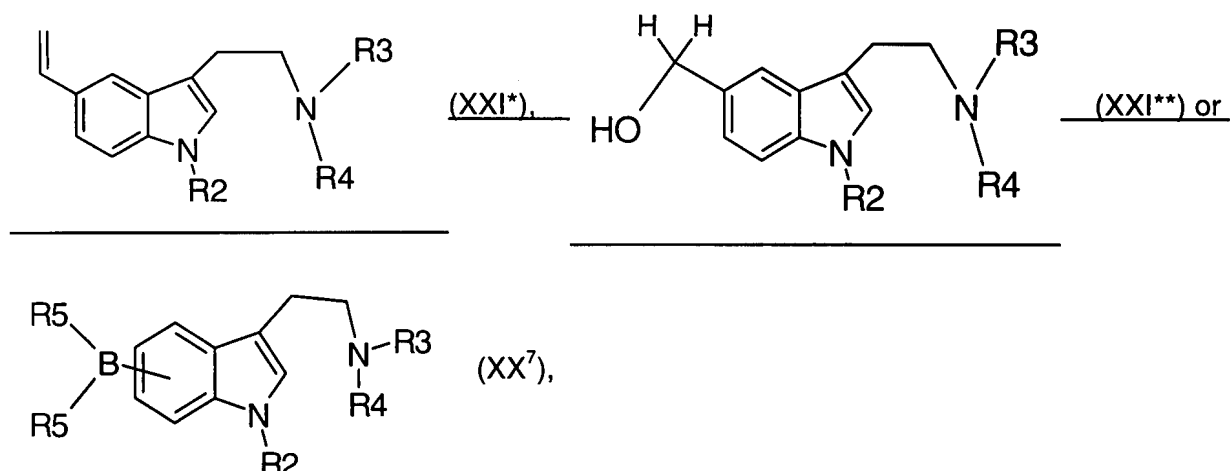
wherein each of R<sub>5</sub> and R<sub>6</sub> independently is an alkoxy residue of a lower alcohol, or the 2 residues R<sub>5</sub> together are C<sub>2</sub>-C<sub>8</sub>alkylene-dioxy, and subsequent hydrolysis, to obtain the compound of the formula XX<sup>7</sup>.



36. (currently amended): A compound of the formulae XIIIa, XVI, II<sup>1</sup>, ~~XII<sup>1</sup>~~, ~~XIV<sup>1</sup>~~, II<sup>2</sup>, XII<sup>2</sup>, XIV<sup>2</sup>, II<sup>3</sup>, XII<sup>3</sup>, XIV<sup>3</sup>, II<sup>4</sup>, XII<sup>4</sup>, XIV<sup>4</sup>, II<sup>5</sup>, XII<sup>5</sup>, XX<sup>7</sup>, XXI\* or XXI\*\*







wherein

n is a number from 0 to 4,

each R<sub>1</sub> is, independently of the other substituents R<sub>1</sub>, unsubstituted or substituted alkyl, unsubstituted or substituted aryl, unsubstituted or substituted heterocyclyl, alkylsulfonyl, sulfonyl alkyl, N-mono- or N,N-disubstituted or unsubstituted aminosulfonyl alkyl, hydroxy, mercapto, nitro, halogen, cyano, carboxamido, N-mono- or N,N-disubstituted carboxamido, carboxhydrazido, unsubstituted or substituted alkoxycarbonyl, unsubstituted or substituted alkoxy, formyl or other alkanoyl, unsubstituted or substituted alkenyl, unsubstituted or substituted alkynyl, unsubstituted or substituted cycloalkyl, alkanoyloxy, N-mono- or N,N-disubstituted or unsubstituted amino, unsubstituted or substituted hydrazino, or is a residue of a boronic acid or an ester thereof;

R<sub>2</sub> is hydrogen or unsubstituted or substituted alkyl, unsubstituted or substituted alkoxycarbonyl, unsubstituted or substituted arylsulfonyl, unsubstituted or substituted alkylsulfonyl, unsubstituted or substituted aryl, carbamoyl or N-mono- or N,N-disubstituted carbamoyl, silyl substituted by three moieties independently selected from unsubstituted or substituted alkyl and substituted or unsubstituted aryl, or acyl, and

R<sub>3</sub> and R<sub>4</sub> are, independently of each other, unsubstituted or substituted alkyl or together form an unsubstituted or substituted alkylene bridge (thus forming a ring with the binding nitrogen) or an alkylene bridge to which a phenyl or a C<sub>3</sub>-C<sub>8</sub>-cycloalkyl ring is condensed at two vicinal carbon atoms of the alkylene bridge,

Hal is nitro or halogen,

Z is unsubstituted or substituted alkyl,

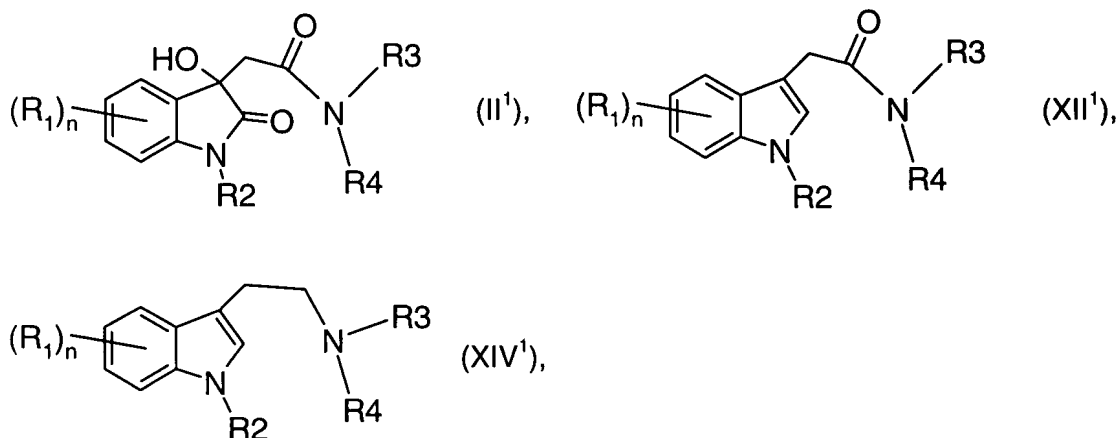
Z\* is unsubstituted or substituted alkyl, unsubstituted or substituted aryl, unsubstituted or substituted arylsulfonyl, unsubstituted or substituted alkylsulfonyl, (Y)<sub>2</sub>N-sulfonyl wherein each Y, independently of

the other, is hydrogen or unsubstituted or substituted alkyl; or Z\* is alkoxy carbonyl, cyano or unsubstituted or substituted heterocyclyl, and

R<sub>5</sub> is unsubstituted or substituted alkyl, or unsubstituted or substituted aryl, with the proviso that n is 1 or 2 in compounds of the formula II<sup>+</sup>, XII<sup>+</sup> and XIV<sup>+</sup>, or a salt thereof.

37-40. (cancelled).

41. (new) A compound of the formulae II<sup>1</sup>, XII<sup>1</sup>, or XIV<sup>1</sup>



wherein

n is 1 or 2,

each R<sub>1</sub> is, independently of the other substituents R<sub>1</sub>, unsubstituted or substituted aryl or unsubstituted or substituted heterocyclyl;

R<sub>2</sub> is hydrogen or unsubstituted or substituted alkyl, unsubstituted or substituted alkoxy carbonyl, unsubstituted or substituted arylsulfonyl, unsubstituted or substituted alkylsulfonyl, unsubstituted or substituted aryl, carbamoyl or N-mono- or N,N-disubstituted carbamoyl, silyl substituted by three moieties independently selected from unsubstituted or substituted alkyl and substituted or unsubstituted aryl, or acyl, and

R<sub>3</sub> and R<sub>4</sub> are, independently of each other, unsubstituted or substituted alkyl or together form an unsubstituted or substituted alkylene bridge (thus forming a ring with the binding nitrogen) or an alkylene bridge to which a phenyl or a C<sub>3</sub>-C<sub>8</sub>-cycloalkyl ring is condensed at two vicinal carbon atoms of the alkylene bridge, or a salt thereof.